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				Date of Report PRISCA	28/5/2022 5.1.0.17
Patient Data					011017
Name	Μ	IRS. RASHMI	Patient ID		012205260144
Birthday		25-1-1983	Sample ID		11304279
Age at term		39.09	Sample Date		26/5/2022
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	13+2
PAPP-A	5.6 mIU/ml	0.89	Method		CRL (<>Robinson)
fb-hCG	27.5 ng/ml	0.65	Scan date		25/05/2022
Risks at sampling date			Nuchal Translucency 1		
Age Risk	ge Risk 1:98		Nuchal Translucency MoM 0.57		
Biochemical T21 risk	mical T21 risk 1:1201		Nasal Bone Present		
Combined T21 risk		1:5788			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1: 00 1:250 Cut off			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 5788 women with the same data, there is one woman with a trisomy 21 pregnancy and 5787 women with not affected pregnancies.		
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18+NT			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		
Risl	x Above Cut Off		Risk above Ag	e Risk	Risk below Age risk